REMARKS

The specification has been amended, as in the parent case, to include identification of sequences. The sequence listing submitted herewith corresponds with that submitted in the parent case.

The applicants intend to rely on the computer readable format (CRF) of the sequence listing as filed in the parent case. The sequence listing submitted herewith does not include new matter and the listing and CRF are the same.

The present divisional application is directed to subject matter that was non-elected in the applicants' parent case.

A PTO-1449 listing the art of record in the parent case is attached.

Respectfully submitted,

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APPENDIX

Version with Markings to Show Changes Made

IN THE SPECIFICATION

Page 16, 3rd ¶ of Example 3, line 27, has been changed to read as follows:

Two carbon probe electrodes were placed into the sample and 4-8 V (d.c.) was applied (power supply; Thurlby 30V, 2A) for between 0.5 to 2 minutes. The cell debris was pelleted and supernatants were analysed by PCR. PCR conditions were as follows; 0.1 µl/ml of sample in PCR buffer (as above), 1 µM (each) of primers ATGCGTCCGGCCGTAGAGGAT SEQ ID No. 1 and GTATCACGAGGCCCTT SEQ ID No. 2, 200 µM of each of dATP, dCTP, dGTP, dTTP, 5U/ml AmpliTaq DNA polymerase (Perkin Elmer). All reagent concentrations are given as the final concentration in a reaction volume made up with PCR buffer (as above). Amplified DNA was analysed on agarose gels stained with ethidium bromide. An amplified DNA fragment of the expected molecular weight (417 bp) was observed in samples which had been subjected to the shortest test time of 30 seconds (see Figure 3). The density of the bands indicated that cell lysis, induced by an applied voltage, released DNA in excess of the background (non-lysed cells control) level.

Attached is paper copy of the Sequence Listing, numbered pages 1-2.